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(54) Process for producing porous polyolefin films

(57) The process comprises blending 100 parts by weight of a polyolefin resin with 50 to 500 parts by weight of barium sulfate preferably having an average particle diameter of 0.1 to 7 μ m, melting the resulting resin composition and forming it into a film, and then stretching the film at least uniaxially by a factor of 1.5 to 7. The polyolefin resin preferably has a melt index of 0.5 to 7 and a density of 0.915 to 0.935; more preferably, it is a linear low-density resin having a melt index of 0.5 to 8.5 and a density of 0.915 to 0.935. As the linear low-density polyethylene resin, a copolymer of ethylene and hexene and/or octene is particularly preferred.

SPECIFICATION

Process for producing porous films

		_
5	Background of the Invention a) Field of the Invention	5
	This invention relates to an improved process for <u>producing porous films</u> . More particularly, it relates to a <u>process for producing porous films</u> which comprises <u>blending a polyolefin resin with barium sulfate as a filler</u> , melting the resulting resin composition and forming it into a film, and then stretching the film at least <u>uniaxially</u> .	10
15	b) Description of the Prior Art Conventionally, there are a large number of well-known processes for producing porous films by forming a resin composition comprising a polyolefin resin and any of various noncompatible fillers into a film and then stretching this film. For example, Japanese Patent Laid-Open Nos. 47334/'82 and 203520/'82 disclose a process for producing porous films which comprises melting a resin composition obtained by blending a polyolefin resin with a filler and liquid	15
20	rubber or a hydroxylated poly-saturated-hydrocarbon, forming the molten room a sheet or film, and then stretching this sheet or film. Moreover, Japanese Patent Laid-Open No. 15538/'83 discloses a process for producing porous films which comprises melting a resin composition obtained by blending a linear low-density polyethylene resin with a filler and a composition obtained by blending a linear low-density polyethylene resin composition into a sheet or film,	20
25	and then stretching this sheet or film. However, the films produced by these processes are disadvantageous in that they exhibit surface tackiness due to the aforesaid component contained in addition to the polyolefin resin and the filler and in that they can only be practically used in relatively large thickness because of their low mechanical strength. Furthermore, it is described in Japanese Patent Laid-Open No. 149303/'83 that such porous films can be used as a leakproof sheet in disposable diapers. Porous films for use as the	25
30	leakproof sheet of a disposable diaper are produced by blending 100 parts by weight of a polyolefin resin with 28 to 200 parts by weight of a filler and 10 to 70 parts by weight of a liquid or waxy hydrocarbon polymer, forming the resulting resin composition into a film, and liquid or waxy hydrocarbon polymer, forming the resulting resin composition into a film, and liquid or waxy hydrocarbon polymer, forming the resulting resin composition into a film, and	30
35	for producing porous films has the disadvantages that some types of files give post stretchability and hence fail to provide fully uniform pores and that the resulting film tends to produce a disagreeable noise. Moreover, the concurrent use of a hydrocarbon polymer makes this process unsafetatory because the hydrocarbon polymer tends to bloom to the film surface	35
40	Such porous films are also useful as a leakproof sheet in sanitary napkins. Conventionally, a sheet of paper which has been rendered liquid-impermeable by means of a synthetic resin such as polyethylene has been used for this purpose. However, the resulting sanitary napkins have the disadvantage of causing a disagreeable sensation during prolonged use because of their lack of permeability to water vapor.	40
45	A sala a a a a sala di la constanti di la cons	45
50	It is another object of the present invention to provide a process for producing porous must which have sufficiently high porosity and thereby exhibit high moisture permeability and gas permeability while retaining excellent water resistance. It is still another object of the present invention to provide a process for producing porous films which are free from surface tackiness, have excellent softness and show little reduction in	50
55	strength. It is a further object of the present invention to provide an improved leakproof sheet for use in disposable diapers. It is a further object of the present invention to provide an improved leakproof sheet for use in	55
	sanitary napkins. Other objects of the present invention will be appar in from the following description. According to the present invention, there is provided a process for producing porous films which comprises melting a resin composition consisting is so ntially of 100 parts by weight if a which comprise and 50 to 500 parts by weight of barium sulfate, forming the molten resin	60
	opolyof the resin and 50 to 500 parts by weight of 2. composition into a film, and then stretching the film at 1 ast uniaxially by a factor of 1.5 to 7. The expression "a resin composition consisting essentially of 100 parts by weight of a polyolefin resin and 50 to 500 parts by weight of barium sulfate" as used herein means that the resin composition may further contain (1) at least one additive selected from common stabilizers, antioxidants, colorants, ultraviolet light absorbents and laubricants and recommons.	65

	addition to the barium sulfate, other in rganic fillers (such as calcium carbonate and the like) or common inorganic and organic m difiers in an amount less than that of barium sulfate used (f r example, not greater than 20% based on the amount of barium sulfate used), but the addition of liquid rubber, a hydroxylated poly-saturated-hydrocarbon or a hydrocarbon polymer as described in the aforementioned Japanese Patent Laid-Open Nos. 47334/'82, 203520/'82, 15538/'83 and 149303/'83 should be positively avoided in order to obtain a porous film free from surface tackiness.	<i>₩</i> 5
10	According to the present invention, porous films which are free from surface tackiness and have excellent properties and which have been unobtainable in the prior art can be produced without using any of the above-described additives used in the prior art. This can be accomplished simply by specifying the type of the filler, its amount used and preferably its average particle diameter: preferably using a low-density polyethylene resin having a specific	10
15	melt index and density and more preferably a linear low-density polyethylene resin (in particular, a linear low-density polyethylene resin comprising a copolymer of ethylene and hexene and/or octene) which most preferably has a specific melt index and density; and stretching the film by a specific factor.	15
20	Detailed Description of the Invention The polyolefin resins which can be used in the present invention include homopoloymers such as polypropylene, low-density polyethylene, high-density polyethylene, linear low-density polyethylene, polybutylene, etc.; copolymers such as ethylene-propylene copolymer, ethylene-butylene copolymer, ethylene-vinyl acetate copolymer, etc.; and blends thereof. Among these resins, low-density polyethylene resins having a melt index of 0.5 to 7 (0.5 to	20
25	8.5 for linear low-density polyethylene resins) and a density of 0.915 to 0.935 are preferred. Although it is preferable to use a single resin, any blend of polyethylene resins having different densities may be used. More preferably, the melt index is in the range of 1 to 5. If the melt index is less than 0.5 or greater than 7 (8.5 for linear low-density polyethylene resins), the resin will have significantly decreased formability into a film and may fail to yield a film of small and	25
30	uniform thickness. On the other hand, if the density is lower than 0.915 or higher than 0.935, the resin will have decreased stretchability and increased rigidity and may fail to yield a soft film. Particularly preferred polyolefin resins are linear low-density polyethylene resins. Linear low-density polyethylene resins are copolymers of ethylene and one or more α-olefins, and differ	30
35	from low-density polyethylene resins prepared by the conventional high-pressure process. Linear low-density polyethylene resins are prepared by the low-pressure process, and useful α-olefins include butene, hexene, octene, decene and the like. The difference between high-pressure-processed low-density polyethylene resins and low-pressure-processed low-density polyethylene	35
40	resins lies in the fact that, when seen from the viewpoint of chemical structure, the former ones are highly branched polymers while the latter ones are straight-chain polymers. Among such lienar low-density polyethylene resins, copolymers of ethylene and hexene and/or octene are particularly preferred.	40
45	In the practice of the present invention, it is preferable to use barium sulfate having an average particle diameter of 0.1 to 7 μ m and more preferably 0.5 to 5 μ m. If the average particle diameter is smaller than 0.1 μ m, well-defined pores cannot be obtained. On the contrary, if it is larger than 7 μ m, the resulting film will have poor stretchability and, therefore, well-defined pores can hardly be obtained just as in cases where the average particle diameter is too small.	45
50	The <u>barium sulfate is used in an amount of 50 to 500 parts</u> by weight, and preferably 100 to 400 parts by weight, per 100 parts by weight of the polyolefin resin. If the amount of barium sulfate used is less than 50 parts by weight, sufficiently high porosity cannot be obtained, while if it is greater than 500 parts by weight, the resulting film cannot be fully stretched because of its increased rigidity and, therefore, will show a reduction in porosity.	50
55	It is preferable to subject the barium sulfate to surface treatment with a fatty acid or a metallic salt thereof, silicone, silane, a resin acid or the like, because this treatment is effective in improving its dispersibility in the resin and producing well-defined pores. So far as the effects of the present invention are not impaired, other inorganic fillers such as calcium carbonate and the like or common inorganic and organic modifiers may be used in addition to the barium sulfate. However, these additives should be used in an amount of not	55
60	greater than 20% based on the amount of barium sulfate us d. Now, the present process for producing porous films will be specifically described hereinbe-	60
65	low. According to the node, at least in additive selected from stabilizers, antioxidants, colorants, ultraviolet light absorbents and lubricants is added to a polyolefin resin and barium sulfate. These ingredients are mix diwith a Hinschel mixer, super-mixer or tumbling mixer. The reafter, using an irdinary single-screw in twin-screw extruder, the resulting mixture is blended and	65

5	pelletized. Then, using an inflation xtruder r T-di xtrud r, these pell ts (al ne r in admixture with polyolefin resin pellets) are melted at a temp rature higher than the melting point of the polyolefin resin (preferably, by 20°C or more) and lower than the decomposition temperature thereof, and formed into a film. In some cases, the aforesaid mixture may be directly formed into a film with an extruder, instead of being pelletized. Subsequently, the film is at east unaxially stretched by a factor of 1.5 to 7 according to a conventional technique such as	5
10	more directions. In the case of biaxial stretching, however, it is preferable to stretch the film simultaneously in the two directions. In order to enhance the morphological stability of pores, the stretched film may be annealed by heating. The porosity is determined by the amount of barium sulfate used, the stretching factor and the like. If the stretching factor is less than 1.5, sufficiently high porosity cannot be obtained,	10
15	while if it is greater than 7, a porous film cannot be steadily produced because of its frequent breakage during the stretching process. Porous films produced by the process of the present invention are characterized by high porosity, excellent softness and little reduction in strength. Moreover, since the good affinity between the polyolefin resin and barium sulfate provides	15
20	good stretchability, not only good workability but also an even distribution of poles can be achieved and, therefore, a porous film can be produced steadily. Especially when a linear low-density polyethylene resin is used as the base resin, the resulting film will show very little and testing in strength. Thus, it is possible to produce porous films which are thinner (for	20
25	example, approximately 10 µm in thickness) than those produced by the prior-art processes. Furthermore, since the resin composition does not contain any of the liquid rubber, hydroxylated poly-saturated-hydrocarbons and hydrocarbon polymers used in the prior-art processes, the resulting porous film is free from surface tackiness. Thus, the porous films of the present invention have sufficiently high porosity and hence	25
30	exhibit good moisture permeability and gas permeability while retaining excellent water resistance, so that they can be used in clothing and sanitary applications. In addition, they can also be used as a filtering medium owing to their even distribution of pores. The properties is further illustrated by the following examples. However, these examples	30
	are given for purposes of illustration only and are not to be construed to limit the scope of the invention. In the examples, melt index (MI) was determined according to ASTM D-1238 and density was determined according to ASTM D-1505.	35
35	(1) Strength and elongation Using a Tensilon tensile testing machine, a piece of film measuring 25 mm (wide) × 100 mm (long) is tested at a extraining rate of 200 mm/min. Its strength and elongation at breakage are	40
40	determined with respect to the machine direction (MD) and the transverse direction (TD). (2) Moisture permeability Moisture permeability is tested according to ASTM E96 (Method D).	40
45	(3) Softness Softness is evaluated by the feel and rated according to the following cristeria:	45
50	-	50
55	Examples 1-19 and Comparative Examples 1-9 Each of the fillers given in Table 1 was added to the corresponding base resin in the amount given in Table 1, and mixed therewith by means of a Henschel mixer. Thereafter, using a twinscrew mixer, the resulting mixture was intimately blended and formed into pellets. Then, using a T-die extruder, these pellets were melted at a temperature 80°C higher than the melting point of the base resin and formed into a film. This film was uniaxially or biaxially (Example 3) stretched by the factor given in Table 1 to obtain a porous film having the thickness given in Table 1.	55
60	However, the film was not stretched in Comparative Example 1 and could not be stretched into a porous film in C mparative Examples 2 and 9. In Comparative Examples 5 and 8, the film could only be stretched by a factor of up to 2. In Comparative Examples 3 and 7, no sampling was possible because of frequent breakage during the stretching process. The strength elongation moisture permeability and softness of the porous films thus obtain described in Comparative Example 1 and could not be stretched in Comparative Example 1 and could not be stretched in Comparative Example 1 and could not be stretched in Comparative Example 1 and could not be stretched in Comparative Example 1 and could not be stretched in Comparative Example 1 and could not be stretched in Comparative Examples 5 and 8, the film could only be stretched in Comparative Examples 2 and 9. In Comparative Examples 5 and 8, the film could only be stretched by a factor of up to 2. In Comparative Examples 3 and 7, no sampling was possible because of frequent breakage during the stretching process.	60
65	were evaluated according to the above-described procedures and the results are shown in Tabl 1.	65

Table l

				Base resin		
			Type ¹⁾	Trade name (manufacturer)	Melt index (g/10 min)	Density (g/cm ³)
)	Example	1	LDPE	REXLON F-41 (Nippon Petrochemicals Co., Ltd.)	5.0	0.923
5	:1	2	•	19	Ħ	67
	19	3	eı	11	**	**
5	tr	4	•	MIRASON 45 (Mitsui Polychemicals Co., Ltd.)	1.5	0.920
	11	5	11	ti	n	
5	ţŝ	6	is .	UBE POLYETHYLENE F0191 (Ube Kosan K.K.)	0.9	0.912
)	n	7	19	Neo-zex 4330 (Mitsui Petrochemical Indus- tries, Inc.)/UBE POLYETHYLENE VF430 (Ube Kosan K.K.)=2/1	3.0	0.940
5	•1	8	L-LDPE	NUCG-5511 (Nippon Unicar Co., Ltd.)	1.0	0.920
	*1	9		11	er	17
)	91	10	**	Ultzex 2020L (Mitsui Petrochemical Indus- tries, Inc.)	2.1	"
=	**	11	••	11	ti ·	"
5	u	12	"	11	м	11
		13	y	u	.**	17

Table 1 (Cont'd)

5		Filler		н		5
	Туре	Average particle dia- meter (µm)	2) Amount (phr)	Stretching factor	Film thickness (µm)	10
10	BaSO ₄	0.8	150	4	40	
15	71	91	**	5	91	15
	81	11	Ħ	2 X 2	tr	
20	11	1.2	130	5	50	20
	81	5.0	120	" .	**	_
25	**	1.2	et .	3 ³)	**	25
30	er .	∵ n	100	4	tr	30
35	**	51	99	**	30	35
	**	88	150	5	15	
40	. ••	4.2	50	6.5	40	40
45	••	0.8	150	5	•	45
	••	0.5	100	7	10	
	**	"	300	3	40	
50			<u></u>	······································		50

Table 1 (Cont'd)

			,			- ,	
5	Stren (kg/25	gth mm)	Elongation (%)		Moisture permeability	Soft-	5
	MD	TD	MD	TD	(g/m ² /24 hr)	ness	
10	6.0	1.5	80	350	3,500	A	10
15	6.5	1.3	60	300	4,200	Λ	15
	4.9	4.5	200	200	3,900	A	
20	5.5	1.5	120	450	3,000 :	. А	20
	5.0	1.3	100	390	62	Α	
25	3.9	1.0	55	390	1,100	В	25
30	4.5	*	50	200	2,000	С	30
35	3.5	1.2	100	420	3,900	С	35
	2.5	0.8	80	400	5,000	Л	
40	7.8	1.4	65	170	2,800	λ	40
45	6.5	1.8	130	510	7,500	А	45
40	2.3	0.7	40	120	8,500	λ	. •
	5.3	1.5	78	360	5,500	Λ	

Table 1 (Cont'd)

5			Base resin						
			Type ¹⁾	Trade name (manufacturer)	Melt index (g/10 min)	Density (g/cm ³)	_		
10	Example	14	L-LDPE	Ultzex 3010F (Mitsui Petrochemical Indus- tries, Inc.)	1.3	0.930			
15	11	15	**	Ultzex 2020L (Mitsui Petrochemical Indus- tries, Inc.)	2.1	0.920	_		
20	41	16	ft	Ultzex 20100J (Mitsui Petrochemical Industries, Inc.)	8.0	0.920	_		
25	#1	17	HDPE	Hi-zex HZ5000S (Mitsui Petrochemical Industries, Inc.)	0.9	0.954	ā		
	11	18	PP	MITSUI NOBLEN JS-G (Mitsui Toatsu Chemicals, Inc.)	1.5	0.890	-		
30	91	19	EPC	MITSUI NOBLEN MJS-G (Mitsui Toatsu Chemicals, Inc.)	"	u			

Table 1 (Cont'd)

5		Fill	er							5
10	Туре	Averac particle meter ()	dia-	2) Amount (Phr)		Str f	etching actor	Film thickne (µm)	ess	10
10 -	BaSO ₄	5.0]	L50		5	50		
15	to .	0.5		9	500		2	40		15
20	n	0.8		2	200		3	t 9		20
25	"	11 11		נ	L50		5	61		25
30 .	"	n			ft		tı .	11		30
	n	ti			•• ·		\$ 1	87		
35	Stre (kg/2	ength !5 mm)	Elo	ngat	ion (%)			ture bility	Soft-	35
40	MD	TD	MD		TD		(g/m ² /	24 hr)	ness	40
45	5.1	1.4	11	0	400		6,7	00	A	45
	5.0	1.0	3	5	20		5,0	00	A	
50	5.5	1.4	10	5	420		4,7	00 .	А	50
55	8.0	2.2	7	0	400		4,5	00	С	55
60	10.3	2.5	9(0	350		4,9	00	С	60
65	9.1	2.3	8:	В	520		4,0	00	С	65
							,			

Table 1 (Cont'd)

_	,	Base resin						
5			Type ¹⁾	Trade name (manufacturer)	Melt index (g/10 min)	Density (g/cm ³)	10	
10		omparative L-LDPE		Ultzex 2020L (Mitsui Petrochemical Indus- tries, Inc.)	2.1	0.920	10	
15	**	2	11	41	"	tt	15	
		3	••	11	11	11		
20	" 4		71	11	80	11	20	
	n	5	••		45	**		
25	et	6	"	NUCG-5511 (Nippon Unicar Co., Ltd.)	1.0	••	25	
	н	7	**	11	tt	11		
30	41	8	n	11	**	"	30	
35	"	9	LDPE	PETROSEN 207 (Toyo Soda Manufacturing Co., Ltd.	8	0.924	35	
40	Notes: 1) LDPE = low-density polyethylene.							
45		2)	Parts b	by weight of the filler of the base rein.	per 100 pa	rts by	45	
50			can be	7) The maximum value a performed steadily. No sampling was possibling during the stretching	le because o	•	50	

Table 1 (Cont'd)

5		Filler				5
10	Type	Average particle dia- meter (µm)	2) Amount (Phr)	Stretching factor	Film thickness (µm)	
10	BaSO ₄	0.8	150	Unstretch- ed	50	10
15	ti	tt	600	Unstretch- ed	-	15
	F1		150	84)	-	
20	CaCO ₃	1.0	ti	4	70	20
_	Glass beads	1.2	120	2 ⁵)	50	_
25	BaSO ₄	ti	30	5	40	25
_	87	••	100	86)	-	
30 _	Glass beads	11	.,	2 ⁷⁾	60	30
35 _	BaSO ₄	ti	130	Unstretch-	_	 35

Table 1 (Cont'd)

						 _	
5	Strength (kg/25 mm)		Elongation (%)		Moisture permeability	Soft-	5
_	MD	TD	MD	TD	(g/m ² /24 hr)		10
10 -	2.5	2.3	570	480	15	A	
15	_		-	-	-	-	15
-	· ·		<u>-</u>	-	-	-	
20	4.3	0.8	20	100	3,000	С	20
20 _	1.5	0.7	140	290	30	С	
25	4.5	1.0	105	430	15	С	25
•	-	_	-	-	<u>-</u>	-	
30	1.0	0.5	40	100	40	С	30
35	_	-	-	-	. -		35

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Example 24

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The same resin composition as used in Example 22 was formed into a film. Using a roll heated to 50°C, this film was uniaxially stretched by a factor of 4 to obtain a porous film having a thickness of 15 μ m. Properties of this porous film were evaluated and the results are shown in Tabl 2.

On the uter side of this porous film was superposed a LDPE film of 70 µm thickn as having apertures of 1 mm diameter (20/cm²) all over its surfact. Disposable diapers using this composite material as the I also exhibited as good performance as those of Examples 20–23.

Table 2

5			Base	e resin	Fi	.ller		5
			Туре	Amount (parts by weight)	Type	Amount (parts by weight)		10
10	Example	20	LDPE	100	BaSO ₄	150		
	11	21	11	н	61	41		
15	81	22	L-LDPE	10	"	200		15
	n	23	LDPE	n	"	150		
20	11	24	L-LDPE	11	11	200		20
	Comparative Example 10		11	15	"	150		
25	"	11	11	99	11	600		25
	"	12	"	88	, 10	150		
30	61	13	**	11	CaCO ₃	40	1_	30
	"	14	**		Glass beads	120	<u> </u>	35
35	L		1					

^{*)} An attempt was made to stretch the film by a factor of 8, but no sampling was possible because of frequent breakage during the stretching process.

^{**)} The film could not be stretched by a factor of more than 2.

Table 2 (Cont'd)

5 10 ·	Stretching factor	Strength [MD/TD] (kg/25 mm)	Moisture permeability (g/m ² /24 hr)	Softness	5
10	4	6.0/1.5	3,500	λ	10
_	. 5	6.5/1.3	4,200	λ	
15	3	5.5/1.3	4,600	A	15
_	2 X 2	4.9/4.5	3,900	A	
20	4	6.7/0.9	4,700	Α	
	Unstretch- ed	2.3/2.2	15	А	20
25 _	Unstretch- ed	-	-	-	25
	8*)	-	-	-	•
30	. 4	4.3/0.8	3,000	c	30
35	2**)	1.5/0.7	30	С	_
					35

Comparative Example 15 120 parts by weight of calcium carbonate having an average particle diameter of 1.2 μm and 20 parts by weight of a hydroxylated poly-saturated-hydrocarbon (Liquid Polybutadiene GI-2000; Nippon Soda Co., Ltd.) were added to 100 parts by weight of linear low-density polyethylene (L-LDPE) having a melt index (MI) of 5, and mixed therewith by means of a 5 Henschel mixer. Thereafter, using a twin-screw mixer, the resulting mixture was intimately blended and formed into pellets. Then, using a 40 mm inflation extruder, these pellets were formed into a film. This film was roll stretched at 80°C by a factor of 3.0 to obtain a porous film having a thickness of 50 μm . This porous film varied in moisture permeability according to the 10 location and exhibited a slight degree of surface tackiness. Disposable diapers using this porous 10 film as the leakproof sheet caused a slight rash on the skin of infants. Comparative Example 16 The procedure of Comparative Example 15 was repeated except that liquid polybutadiene 15 (Nisso PBG; Nippon Soda Co., Ltd.) or rubbery EPR (Toughmer P0480; Mitsui Petrochemical 15 Industries, Inc.) was used as the hydroxylated poly-saturated-hydrocarbon. Thus, there were obtained films having a thickness of 50 μm . These porous films exhibited surface tackiness and varied in moisture permeability according to the location. Disposable diapers using each of these porous films as the leakproof sheet caused a slight rash on the skin of infants. Porous films obtained by the process of the present invention have high porosity and excellent 20 softness and, moreover, show little reduction in strength in spite of their small thickness, so that they are very suitable for use as a leakproof sheet in sanitary napkins. Conventional sanitary napkins are so constructed that a liquid absorber such as fluffy pulp, cotton, absorbent resin or the like is partially covered with a film of paper having been rendered liquid-impermeable by 25 treatment with a synthetic resin such as polyethylene or the like and the resulting structure is 25 then wrapped in a non-woven fabric. In sanitary napkins using the porous film of the present invention as the leakproof sheet, this leakproof sheet has a large number of pores which allow water vapor to pass therethough. Accordingly, they can keep the skin of the user in a dry state and cayse no disagreeable sensation even during prolonged use. The following examples illustrate the use of the porous film of the prsent invention as a 30 leakproof sheet in sanitary napkins. Examples 25-27 Barium sulfate having an average particle diameter of 0.8 μm was added to 100 parts by 35 weight of linear low-density polyethylene (L-LDPE) having a melt index (MI)) of 2.1 in the 35 amount given in Table 3, and mixed therewith by means of a Henschel mixer. Thereafter, using a twin-screw mixer, the resulting mixture was intimately blended and formed into pellets. Then, using a T-die extruder, these pellets were melted at 230°C and formed into a film. This film was uniaxially stretched between a preheating roll heated to 80°C and a stretching roll by the factor 40 given in Table 3 to obtain a porous film having a thickness of 20 μm. Properties of this porous 40 film were evaluated and the results are shown in Table 3. Sanitary napkins were made by covering a filling of fluffy pulp partially with each of the porous films obtained in Examples 25-27, wrapping the resulting structure in a non-woven fabric and then heat sealing its overlapping portions. When these sanitary napkins and commercially available ones having a 45 liquid-impermeable film of polyethylene-coated paper were comparatively tested by using them 45 practically for prolonged periods of time, the sanitary napkins in accordance with the present

invention did not cause a disagreeable, stuffy sensation.

/		
/	9	
	Table	
	HI	\iint
/		

	Bas	Base resin	E4.	Filler				
	Ē	Amount		Amount	Stretching	[MD/TD]	Moisture permeability Soft-	Søft-
	1 y pe	(parts by 1ype weight)	lype	(parts by weight)	4	(kg/25 mm)	(kg/25 mm) (g/m ² /24 hr)	กеรธ
Example 25	L-LDPE	100	BaSO4	100	9	3.2/1.0	5,200	A
Example 26	=	=	=	150	Ŋ	3.0/0.9	4,800	∢
Example 27	=	=	=	400	m	2.5/0.8	4,200	~

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5	CLAIMS 1. In a process for producing porous films which comprises melting a resin composition consisting essentially of a polyolefin resin and a filler non-compatible therewith, forming the molten resin composition into a film, and then stretching the film at least uniaxially, the molten resin composition into a film, and then stretching the film at least uniaxially, the improvement in which the non-compatible filler is barium sulfate, the barium sulfate is used in an amount of 50 to 500 parts by weight per 100 parts by weight of the polyolefin resin, and the film is stretched by a factor of 1.5 to 7.	5
	2. A process as claimed in claim 1 wherein the average particle diameter of the barium sulfate is in the range of 0.1 to 7 μm.	10
	sulfate is in the range of 0.5 to 5 µm. 4. A process as claimed in claim 1 wherein the barium sulfate is used in an amount of 100 to 400 parts by weight per 100 parts by weight of the polyolefin resin. 5. A process as claimed in claim 1 wherein the barium sulfate has been subjected to surface treatment with a fatty acid or a metallic salt thereof, silicone, silane or a resin acid. 6. A process as claimed in claim 1 wherein the polyolefin resin is a polyethylene resin.	15
	polyethylene resin having a melt index of 0.3 to 7 and 5 density polyethylene resin is a single 8. A process as claimed in claim 7 wherein the low-density polyethylene resin is a single	20
20	resin. A process as claimed in claim 7 wherein the low-density polyethylene resin is a blend of	
	polyethylene resins having different densities. 10. A process as claimed in claim 7 wherein the melt index is in the range of 1 to 5. 11. A process as claimed in claim 6 wherein the polyethylene resin is a linear low-density	25
25	polyethylene resin. 12. A process as claimed in claim 11 wherein the linear low-density polyethylene resin has a melt index of 0.5 to 8.5 and a density of 0.915 to 0.935. 13. A process as claimed in claim 11 wherein the linear low-density polyethylene resin is a 13.	
	copolymer of ethylene and nexerie ancietopes, moisture permeability and gas permeability which	30
30	has been produced by a process as claimed if any other in disposable diapers. 15. The use of the porous film of claim 14 as a leakproof sheet in sanitary napkins. 16. The use of the porous film of claim 14 as a leakproof sheet in sanitary napkins. 17. The process for producing porous films substantially as hereinbefore described with	35
35	17. The process for producing examples of the foregoing examples as hereinbefore described with reference to any one of the use of a porous film substantially as hereinbefore described with reference to any one of the foregoing examples of the invention.	

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